

# CONTAMAC

510(k) Premarket Notification



## 510 (k) SUMMARY OF SAFETY AND EFFECTIVENESS

This summary of 510(k) safety and effectiveness information is being submitted in accordance with the requirements of SMDA 1990 and 21 CFR 807.92.

The assigned 510(k) number is:

**K023349**

### Applicant information:

Date Prepared:	October 2, 2002
Name:	<b>CONTAMAC Ltd.</b>
Address	Bearwalden Business Park Saffron Walden Essex England CB11 4JX
Contact Person:	Robert McGregor
Phone number:	44-1799 542 000
US Agent:	Medvice Consulting, Inc. Martin Dalsing
Phone number	(970) 243-5490
Fax number	(970) 243-5501

### Device Information:

Device Classification:	Class II
Classification Number:	LPL
Classification Name:	Lenses, Soft Contact, Daily Wear
Trade Name:	<b>CONTAFLEX GM3 58% (acofilcon A) Spherical Soft Contact Lens for Daily Wear (clear and tinted, lathe-cut)</b>

# CONTAMAC

510(k) Premarket Notification



## Equivalent Devices:

The CONTAFLEX GM3 58% (acofilcon A) Spherical Soft Contact Lenses are substantially equivalent to the following predicate device

*Predicate device:* "BENZ 55G" manufactured/distributed by Benz Research and Development.  
510(k) number; K952620

## Device Description:

K018349

The CONTAFLEX GM3 58% Spherical Soft Contact Lenses are fabricated from acofilcon A, which in the dry (unhydrated) state may be machined and polished. The hydrophilic nature of this material allows the lens to become soft and pliable when immersed in an aqueous solution.

The non-ionic lens material, (acofilcon A) is a terpolymer based on high purity Glycerol Methacrylate 2,3-Dihydroxypropyl Methacrylate (GMA), with N-vinyl-2-pyrrolidone (NVP), methyl methacrylate (MMA), and 2-hydroxyethyl methacrylate (2-HEMA) and cross-linked with Diallyl Maleate (DAM). It consists of 42% acofilcon A and 58% water by weight when immersed in normal saline solution buffered with sodium bicarbonate. The lens is available in clear and with a blue visibility-handling tint, Color additive 'Reactive Blue 4' 21 CFR part 73.2121. The acofilcon A name has been adopted by the United States Adopted Names Council (USAN).

In the hydrated state, the lens conforms to the curvature of the eye covering the cornea and extending slightly beyond the limbus forming a colorless, transparent optical surface. The (acofilcon A) soft hydrophilic contact lens has a spherical back surface. The hydrophilic properties of the lens require that it be maintained in a fully hydrated state in a solution compatible with the eye. If the lens dries out, it will become hard and appear somewhat warped however, it will return to its proper configuration when completely rehydrated in the proper storage solution.

The hydrophilic characteristics allow aqueous solutions to enter the lens and in its fully hydrated state the lens is approximately 58% water by weight. The physical properties of the lens are:

<b>Refractive Index</b>	1.52 (dry) 1.40 (hydrated)
<b>Light Transmission</b>	greater than 93%
<b>Water Content</b>	58 %
<b>Specific Gravity</b>	1.103 (hydrated)
<b>Oxygen Permeability</b>	$25.50 \times 10^{-11}$ (cm <sup>2</sup> /sec) (ml O <sub>2</sub> /ml x mm Hg @ 35°C), (revised Fatt method).



## **Intended Use:**

The **CONTAFLEX GM3 58 (acofilcon A) Spherical** Soft Contact Lenses for daily wear are indicated for the correction of visual acuity in aphakic and not aphakic persons with non-diseased eyes with myopia or hyperopia. The lens may be worn by persons who exhibit refractive astigmatism of 0.75 diopters or less where the astigmatism does not interfere with visual acuity. The lens may be disinfected using a chemical disinfecting system.

Eyecare practitioners may prescribe the lenses for frequent/planned replacement wear, with cleaning, disinfection and scheduled replacement. When prescribed for frequent/planned replacement wear, the lens may be cleaned and disinfected using a chemical (not heat) lens care system.

## **Description of Safety:**

A series of preclinical testing were performed to demonstrate the safety and effectiveness of the CONTAFLEX GM3 58 Soft Contact Lens material. The results of all testing demonstrated that the safety and effectiveness of the CONTAFLEX GM3 58 Soft Contact Lens is equivalent to the currently marketed Benz 55G contact lens material. A summary of these results from the preclinical studies is presented below.

### **Toxicology:**

**In-Vitro Cytotoxicity:** ISO 10993-5 was conducted in accordance with standards on test article. The test article meets the requirements of the Agarose Overlay Method.

**Systemic Toxicity:** The lens material meets the requirements of the systemic injection test and is considered non-toxic.

**Acute Ocular Irritation:** Acute ocular irritation test was performed and produced no ocular irritation.

### **Shelf Life**

Shelf life requirements are satisfied via referencing rights granted to Contamac Ltd. for 510(k) 973597. The data presented supports substantial equivalence of this CONTAFLEX GM3 58 Soft Contact Lens material to the already marketed Benz 55G.

### **Solution Compatibility**

Studies were conducted on blue tinted lens material. Lenses were run through 30 cycles of cleaning and conditioning to establish the compatibility of the lens material with the recommended care regimen. The parameters of the base curve, back vertex power, total diameter and overall lens physical appearance were recorded prior to and upon completion of 30 cycles. Initial and final data were compared. There were no significant changes to lens parameters after 30 complete cycles.



---

### **Clinical Testing**

Below is a summary of the clinical study carried out to evaluate the safety and efficacy of the CONTAFLEX GM3 58 Soft Contact Lens material when used as a daily wear contact lens for the correction of visual acuity.

A total of 122 eyes (61 patients) were entered into the study by 5 investigators. Prior to entry into this study each patient was required to read and sign a statement of informed consent. All patients who signed a Statement of Informed Consent are accounted for in this report. Of the 122 eyes (61 patients enrolled), 100 eyes (50 patients) completed the study.

The safety and efficacy measures for the study were:

Safety: Adverse Events, Positive Slit Lamp Findings,  
Symptoms/Complaints and Keratometry Changes.

Efficacy: Refractive Changes, Lens Visual Acuity, Lens  
VA Line Changes, Comfort and Vision, Average Wearing time.

The sponsor concludes that CONTAFLEX GM3 58 Soft Contact Lens material is equivalent in safety and efficacy to the predicate device, the Benz 55G.

### **Substantial Equivalence:**

The CONTAFLEX GM3 58 Soft Contact Lens is substantial equivalence and does not raise different questions of safety and effectiveness than the predicate device identified previously. The difference between the two devices is the USAN name.

The following table depicts the pre-clinical characteristics of the CONTAFLEX GM3 58 material, as well as the predicate device.

# CONTAMAC

510(k) Premarket Notification



**Substantial Equivalence table**

	<b>Pre-Clinical equivalency / Device</b>	<b>Contaflex GM3 58% (acofilcon A)</b>	<b>BENZ 55G (hioxifilcon A)</b>
1.)	<b>Intended Use</b>	Indicated for daily wear for the correction of visual acuity in aphakic and not aphakic persons with non-diseased eyes with myopia or hyperopia.	Indicated for daily wear for the correction of visual acuity in aphakic and not aphakic persons with non-diseased eyes with myopia or hyperopia.
2.)	<b>Functionality</b>	After machining from the optical blank, the contact lenses act as a refractive medium that focus light rays from near and distant objects on the retina.	After machining from the optical blank, the contact lenses act as a refractive medium that focus light rays from near and distant objects on the retina.
3.)	<b>Indications</b>	Daily wear, Soft (hydrophilic) contact lens	Daily wear, Soft (hydrophilic) contact lens
4.)	<b>Production Method</b>	Lathe-cut	Lathe-cut
5.)	<b>FDA Group #</b>	Group # 2 >50% Water, Nonionic Polymers	Group # 2 >50% Water, Nonionic Polymers
6.)	<b>USAN name</b>	Acofilcon A	Hioxifilcon A
7.)	<b>Water Content</b>	58.0%	58.0%
8.)	<b>Oxygen Permeability</b>	25.5 X 10 <sup>-11</sup> (cm <sup>2</sup> /sec) (ml O <sub>2</sub> /ml x mm Hg @ 35 degrees C), (revised Fatt method).	23.0 X 10 <sup>-11</sup> (cm <sup>2</sup> /sec) (ml O <sub>2</sub> /ml x mm Hg @ 35 degrees C), (revised Fatt method).
9.)	<b>Specific Gravity</b>	1.103	1.045



Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

DEC 24 2002

Medvice Consulting, Inc.  
c/o Martin Dalsing  
623 Glacier Drive  
Grand Junction, CO 81503

Re: K023349

Trade/Device Name: Contaflex GM3 58% (acofilcon A) Spherical Soft Contact Lens  
For Daily Wear (clear and tinted, lathe-cut)

Regulation Number: 21 CFR 886.5925

Regulation Name: Soft (hydrophilic) Contact Lens

Regulatory Class: Class II

Product Code: LPL

Dated: October 2, 2002

Received: October 7, 2002

Dear Mr. Dalsing:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

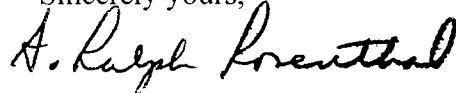
Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set

forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at (301) 594-4692. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address <http://www.fda.gov/cdrh/dsma/dsmamain.html>

Sincerely yours,

A handwritten signature in black ink that reads "A. Ralph Rosenthal". The signature is written in a cursive, flowing style.

A. Ralph Rosenthal, M.D.  
Director  
Division of Ophthalmic and Ear,  
Nose and Throat Devices  
Office of Device Evaluation  
Center for Devices and  
Radiological Health

# CONTAMAC

510(k) Premarket Notification



## INDICATIONS FOR USE STATEMENT

**Device Name:** **CONTAFLEX GM3 58% (acofilcon A) Spherical Soft Contact Lens for Daily Wear (clear and tinted, lathe-cut)**

### **INDICATIONS FOR USE:**

The **CONTAFLEX GM3 58 (acofilcon A) Spherical** Soft Contact Lenses for daily wear are indicated for the correction of visual acuity in aphakic and not aphakic persons with non-diseased eyes with myopia or hyperopia. The lens may be worn by persons who exhibit refractive astigmatism of .75 diopters or less where the astigmatism does not interfere with visual acuity. The lens may be disinfected using a chemical disinfecting system.

Eyecare practitioners may prescribe the lenses for frequent/planned replacement wear, with cleaning, disinfection and scheduled replacement. When prescribed for frequent/planned replacement wear, the lens may be cleaned and disinfected using a chemical (not heat) lens care system.

(PLEASE DO NOT WRITE BELOW THIS LINE - CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence ~~(Division Signature)~~ of Device Evaluation (ODE)

Division of Ophthalmic Ear,  
Nose and Throat Devices

*Daniel W. C. Brown, P.D.*

510(k) Number K023349

Prescription Use X  
(Per 21 CFR 801.109)

or

Over-The-Counter Use     

(Optional Format 1-2-96)